PTO/SB/08A (08-03)
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INFORMATION DISCLOSURE STATEMENT BY APPLICANT

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	Complete if Known
Application Number	10/602,692
Filing Date	June 20, 2003
First Named Inventor	Sommadossi et al.
Group Art Unit	1623
Examiner Name	Travis C. McIntosh, III
Attorney Docket Number	06171.105071 IDX 1006 CON3 US

Sheet 3226087

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			U.	S. PATENT DOCUMENTS			
Examiner Initials *	Cite No. 1		Kind Code (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages/Relevant Figures Appear	T ⁴
Y	AA	6,455,508	B1	Ramasamy et al.	09-24-2002		
	AB	6,495,677	Bl	Ramasamy et al.	12-17-2002		<u> </u>
	AC	6,566,344	Bl	Gosselin et al.	05-20-2003		
	AD	6,566,365	Bl	Storer	05-20-2003		<u> </u>
	AE	6,569,837	Bl	Gosselin et al.	05-27-2003		
	AF	6,605,614	B2	Bachand et al.	08-12-2003		
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	AK	6,812,219	B2	LaColla et al.	11-02-2004		
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13	AAF	2004-0072788	A1	Bhat et al.	04-15-2004		t

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Sheet 3226087

		-	U.S	S. PATENT DOCUMENTS			
Examiner Cite No.		U.S. Patent Doc Number	ument Kind Code (if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages/Relevant Figures Appear	74
Q2	BA	2004-0097461	A1	Sommadossi et al.	05-20-2004		
	BB	2004-0097462	Al	LaColla et al.	05-20-2004		
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	BG	2004-0147464	A1	Roberts, et al.	07-29-2004		
	ВН	2004-0248844	A1	Ismaili <i>et al</i> .	12-09-2004		\Box
	BI	2005-0009737	A1	Clark, et al.	01-13-2005		
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P	ВР	FR	2,662,165	Al	Univ. Pier et Curie	11-22-1991	provided as Derwent Abstract	
0	ВQ	JP	63-215694	A2	Yamasa Shoyu Co. Ltd.	09-08-1988	provided as Delphion Abstract	
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Examiner Signature	Date Considered	1/26/05	
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Sheet 6

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TRADEL MAIN	Complete if Known
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First Named Inventor	Sommadossi et al.
Group Art Unit	1623
Examiner Name	Unassigned
Atterney Docket Number	06171.105071 IDX 1006 CON3

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	U.S. PATENT DOCUMENTS										
Examiner Initials •	Cite No. '		ment Kind Code if known)	Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pgs, Clmns, Lns, Where Relevant Passages/Relevant · Figs Appear					
B	AA	3,480,613	Α	Walton et al.	11-25-1969						
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	AL	2003/028013	Al	Wang et al.	02-06-2003						
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	AN	2003/0060400	Al	LaColla et al.	03-27-2003						
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A	AP	2003/0087873	Al	Stuyver et al.	05-08-2003						

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Examiner Initials •	Cite No. 1		ign Patent Docum Number Kind (if k		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	I.
(A)	AQ	FR	1,521,076	Α	Merck & Co. Inc.	04-12-1968		_
	AR	FR	1,581,628	Α	Merck & Co. Inc.	09-19-1969		
	AS	FR	2,662,165	Α	Univ. Paris Curie	11-22-1991		
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	AV	JP	63-215694	Α	Yamasa Shoyu Co. Ltd.	09-08-1988		
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	AX	wo	98/16184	A2	ICN Pharmaceuticals .	04-23-1998		
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(d)	AAA	wo	01/32153	A2	Biochem Pharma	05-10-2001		

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Examiner	Date	13/2	1/5
Signature	Considered	1 7 8	145

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2000xilute 10	W 1000 1447/0/10			Application Number	10/602,692		
INFO	RMATION	DISCLO	SURE	Filing Date	June 20, 2003		
STATEMENT BY APPLICANT			CANT	First Named Inventor	Sommadossi et al.		
				Group Art Unit	1623		
	(use as many sheets as necessary)			Examiner Name	Unassigned		
Sheet	2	of	6	Attorney Docket Number	06171.105071 IDX 1006 CON3		
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					REIGN PATENT DOCUMENTS			
Examiner Initials •	Cite No.		ign Patent Docum Number Kind (if k		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD- YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	14
(2)	BA	wo	01/60315	A2	Biochem Pharma	08-23-2001		Т
	BB	WO	01/68663	Al	ICN Pharmaceuticals	09-20-2001		П
	BC	WO	01/79246	A2	Pharmasset	10-25-2001		П
	BD	WO	01/90121	A2	Novirio Pharm. (Idenix)	11-29-2001		Γ
	BE	WO	01/91737	A2	Novirio Pharm. (Idenix)	06-12-2001		Г
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	Bl	WO	02/18404	A2	F. Hoffmann-La Roche	03-07-2002		
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	BK	wo	02/48165	A2	Pharmasset	06-20-2002		Г
	BL	wo	02/057287	A2	Merck & Co. Inc.	07-25-2002		Г
	BM	wo	02/057425	A2	Merck & Co. Inc.	07-25-2002		
	BN	wo	02/070533	A2	Pharmasset	09-12-2002		Г
	BO	wo	02/094289	Αl	F. Hoffmann-La Roche	11-28-2002		
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	BQ	WO	03/026589	A2	Idenix; CNRS; U. Montp.	04-03-2003		
	BR	wo	03/026675	Al	Idenix; CNRS; U. Montp.	04-03-2003		Г
	BS	wo	03/051899	AI	Ribapharm	06-26-2003		Г
	BT	wo	03/061385	AI	Ribapharm	07-31-2003		
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Examiner Signature	2	5	Date Considered	3/31/05	5

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6-1	r (przs. 1449A/PTO	WORK REGULERALISM AL		Complete if Known			
Sansurvie 10	rum imanito			Application Number	10/602,692		
INFO	RMATION	DISCLO	SURE	Filing Date	June 20, 2003		
STAT	STATEMENT BY APPLICANT			First Named Inventor	Sommadossi et al.		
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	(use as many she	eu as necessary)		Examiner Name	Unassigned		
Sheet	3	of	6	Attorney Docket Number	06171.105071 IDX 1006 CON3		

3425649 1 OTHER PRIOR ART - NON PATENT LITERATURE DOCUMENTS Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, Examine No. 1 Initials • journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. ALTMANN et al, "The synthesis of 1'-methyl carbocyclic thymidine and its effect on nucleic acid CA duplex stability," Synlett, Thieme Verlag, Stuttgart, De, 10:853-855 (1994). CB BAGINSKI, S. G, et al., "Mechanism of action of a pestivirus antiviral compound," PNAS USA. 97(14):7981-7986 (2000). CC BEIGELMAN, L.N., et al, "Epimerization during the acetolysis of 3-O-acetyl-5-O-benzoyl-1,2-Oisopropylidene-3-C-methyl-α,D-ribofuranose. Synthesis of 3'-C-methylnucleosides with the β-Dribo- and α-D-arabino configurations," Carbohydrate Research, 181:77-88 (1988). BEIGELMAN, L.N., et al, "A general method for synthesis of 3'-C-alkylnucleosides," Nucleic Acids Symp. Ser., 9:115-118 (1981). CE BERENGUER, M., et al, "Hepatitis B and C viruses: Molecular identification and targeted antiviral therapies," Proceedings of the Association of American Physicians, 110(2), 98-112 (1998). CARROLL, S.S., et al., "Inhibition of hepatitis C virus RNA replication by 2'-modified nucleoside analogs," The Journal of Biological Chemistry, 278(14):11979-11984 (2003). CG CZERNECKI, S., et al, "Synthesis of various 3'-branched 2',3'-unsaturated pyrimidine nucleosides as potential anti-HIV agents," J. Org. Chem., 57:7325-7328 (1992). De FRANCESCO, R., et al., "Approaching a new era for hepatitis C virus therapy: Inhibitors of the NS3-4A serine protease and the NS5B RNA-dependent RNA polymerase," Antiviral Research, 58:1-16 (2003). CI FAIVRE-BUET, V., et al, "Synthesis of 1'-deoxypsicofuranosyl-deoxynucleosides as potential anti-HIV agents," Nucleosides & Nucleotides, 11(7):1411-1424 (1992). CJ FARKAS, J., et al., "Nucleic acid components and their analogues. XCIV. Synthesis of 6-amino-9-(1deoxy-β-D-psicofuranosyl)purine", Collect. Czech. Chem. Commun. 32:2663-2667 (1967). CK FARKAS, J., et al., "Nucleic acid components and their analogues. LXXIX. Synthesis of methyl 1deoxy-D-psicofuranosides substituted at C(1) with halo atoms or a mercapto group," Collect. Czech. Chem. Commun., 31:1535-1543 (1996). CL FEDOROV, I.I., et al, "3'-C-Branched 2'-deoxy-5-methyluridines: Synthesis, enzyme inhibition, and antiviral properties," J. Med. Chem., 35(24):4567-4575 (1992). FRANCHETTI, P., et al., "2'-C-Methyl analogues of selective adenosine receptor agonists: synthesis CM and binding studies," J. Med. Chem., 41(10):1708-1715 (1998). GROUILLER, A., et al., "Novel p-toluenesulfonylation and thionocarbonylation of unprotected thymine nucleosides," Synlett, 1993, 221-222 (March 1993). HARAGUCHI, K., et al., "Preparation and reactions of 2'- and 3'- vinyl bromides of uracil nucleosides: Versatile synthons for anti-HIV agents," Tetrahedron Letters, 32(28):3391-3394 (1991).

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Straine in the 144/74110				Application Number	10/602,692		
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STATEMENT BY APPLICANT			CANT	First Named Inventor	Sommadossi et al.		
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Sheet	4	of	6	Attorney Docket Number	06171.105071 IDX 1006 CON3		

3425649 1 OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, xemine Cite No. 1 Initials • journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published. DA HARAGUCHI, K., et al., "Stereoselective synthesis of 1'-C-branched uracil nucleosides from uridine," Nucleosides & Nucleotides, 14(3-5):417-420 (1995). DB HARRY-O'KURU, R.E., et al., "A short, flexible route toward 2'-C-branched ribonucleosides", J.Org. Chem., 62:1754-1759 (1997). (Scheme 11). HARRY-O'KURU, R.E., et al., "2'-C-Alkylribonucleosides: Design, synthesis, and conformation," DC Nucleosides & Nucleotides, 16(7-9):1457-1460 (1997). ["Rogers" in #2; correct name in #7] DD HATTORI, H., et al, "Nucleosides and nucleotides. 175. Structural requirements of the sugar moiety for the antitumor activities of new nucleoside antimetabolites, 1-(3-C-ethynyl-b-D-ribopentofuranosyl)cytosine and -uracil," J. Med. Chem., 41:2892-2902 (1998). HREBABECKY, H., et al., "Nucleic acid components and their analogues. CXLIX. Synthesis of pyrimidine nucleosides derived from 1-deoxy-D-psicose," Collect. Czech. Chem. Commun., 37:2059-2065 (1972). DF HREBABECKY, H., et al. "Synthesis of 7- and 9B-D-psicofuranosylguanine and their 1'-deoxy derivatives," Collect. Czech. Chem. Commun., 39:2115-2123 (1974). DG INO, T., et al., "Nucleosides and nucleotides. 139. Stereoselective synthesis of (2'S)-2'-C-alkyl-2'deoxyuridines," Nucleosides and Nucleotides, 15(1-3):169-181 (1996). DH ITOH, Y., et al, "Divergent and stereocontrolled approach to the synthesis of uracil nucleosides branched at the anomeric position," J. Org. Chem., 60(3):656-662 (1995). DI JOHNSON, C.R., et al, "3'-C-Trifluoromethyl ribonucleosides," Nucleosides & Nucleotides. 14(1&2):185-194 (1995). DJ KAWANA, M., et al., "The deoxygenation of tosylated adenosine derivatives with Grignard reagents," Nucleic Acids Symp. Ser., 17:37-40 (1986). LAVAIRE, S., et al., "3'-Deoxy-3'-C-trifluoromethyl nucleosides: Synthesis and antiviral evaluation," Nucleosides & Nucleotides, 17(12):2267-2280 (1998). LEYSSEN, P. et al., "Perspectives for the treatment of infections with Flaviviridae," Clinical Microbiology Reviews (Washington, D.C.), 13(1):67-82 (January 2000). MARTIN, X., et al., "Intramolecular hydrogen bonding in primary hydroxyl of thymine 1-(1-deoxy-\beta-D-psicofuranosyl) nucleoside," Tetrahedron, 50(22):6689-6694 (1994). DN MATSUDA, A., et al., "Radical deoxygenation of tert-alcohols in 2'-branched-chain sugar pyrimidine nucleosides: Synthesis and antileukemic activity of 2'-deoxy-2'(S)-methylcytidine," Chem. Pharm. Bull., 35(9):3967-3970 (1987). MATSUDA, A., et al., "Alkyl addition reaction of pyrimidine 2'-ketonucleosides: Synthesis of 2'branched-chain sugar pyrimidine nucleosides (Nucleosides and Nucleotides. LXXXI)," Chem. Pharm. Bull., 36(3):945-953 (1988).

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Substitute fo	Substitute for form 1449APTO INFORMATION DISCLOSURE STATEMENT BY APPLICANT	Complete if Known					
amane in				Application Number	10/602,692		
INFO	RMATION	DISCLO	SURE	Filing Date	June 20, 2003		
STAT	STATEMENT BY APPLICANT			First Named Inventor	Sommadossi et al.		
				Group Art Unit	1623		
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	1	OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS	
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	EA	MATSUDA, A., et al., "Nucleosides and Nucleotides. 94. Radical deoxygenation of tert-alcohols in	l
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				Group Art Unit	1623		
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		OTHER PRIOR ART – NON PATENT LITERATURE DOCUMENTS	
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B	FA	SAMANO, V., et al., "Nucleic acid related compounds. 77. 2',3'-Didehydro-2',3'-dideoxy-2'(and 3')-methylnucleosides via [3,3]-sigmatropic rearrangements of 2'(and 3')-methylene-3'(and 2')-O-thiocarbonyl derivatives and radical reduction of a 2'-chloro-3'-methylene analogue," Can. J. Chem., 71:186-191 (1993).	
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